### MEDWATCH
The FDA Safety Information and Adverse Event Reporting Program

**For VOLUNTARY reporting of adverse events, product problems and product use errors**

**Page 1 of 3**

#### A. PATIENT INFORMATION
1. **Patient Identifier**
   - Date of Birth:
2. **Age at Time of Event or Date of Birth:**
3. **Sex**
   - Female
   - Male
4. **Weight**
   - lb
   - kg

#### B. ADVERSE EVENT, PRODUCT PROBLEM OR ERROR
Check all that apply:
1. **Adverse Event**
2. **Product Problem (e.g., defects/malfunctions)**
3. **Product Use Error**
4. **Problem with Different Manufacturer of Same Medicine**

2. **Outcomes Attributed to Adverse Event**
   - (Check all that apply)
   - Death: (mm/dd/yyyy)
   - Disability or Permanent Damage
   - Life-threatening
   - Congenital Anomaly/Birth Defect
   - Hospitalization - initial or prolonged
   - Other Serious (Important Medical Events)
   - Required Intervention to Prevent Permanent Impairment/Damage (Devices)

3. **Date of Event (mm/dd/yyyy)**
4. **Date of this Report (mm/dd/yyyy)**

5. **Describe Event, Problem or Product Use Error**

6. **Relevant Tests/Laboratory Data, Including Dates**

7. **Other Relevant History, Including Preexisting Medical Conditions**
   - (e.g., allergies, race, pregnancy, smoking and alcohol use, liver/kidney problems, etc.)

#### E. SUSPECT MEDICAL DEVICE
1. **Brand Name**
2. **Common Device Name**
3. **Manufacturer Name, City and State**

4. **Model #**
5. **Lot #**

#### C. PRODUCT AVAILABILITY
Product Available for Evaluation? (Do not send product to FDA)
- Yes
- No
- Returned to Manufacturer on: (mm/dd/yyyy)

#### D. SUSPECT PRODUCT(S)
1. **Name, Strength, Manufacturer (from product label)**
   - #1 Name:
   - Strength:
   - Manufacturer:
   - #2 Name:
   - Strength:
   - Manufacturer:

#### F. OTHER (CONCOMITANT) MEDICAL PRODUCTS
Product names and therapy dates (exclude treatment of event)

#### G. REPORTER (See confidentiality section on back)
1. **Name and Address**
   - Name:
   - Address:
   - City: State: ZIP:

2. **Phone #**
3. **E-mail**

4. **Also Reported to:**
   - Manufacturer
   - User Facility
   - Distributor/Importer

5. If you do NOT want your identity disclosed to the manufacturer, place an "X" in this box: [ ]

**FORM FDA 3500 (2/13)** Submission of a report does not constitute an admission that medical personnel or the product caused or contributed to the event.
ADVICE ABOUT VOLUNTARY REPORTING


Report adverse events, product problems or product use errors with:
- Medications (drugs or biologics)
- Medical devices (including in-vitro diagnostics)
- Combination products (medication & medical devices)
- Human cells, tissues, and cellular and tissue-based products
- Special nutritional products (dietary supplements, medical foods, infant formulas)
- Cosmetics
- Food (including beverages and ingredients added to foods)

Report product problems - quality, performance or safety concerns such as:
- Suspected counterfeit product
- Suspected contamination
- Questionable stability
- Defective components
- Poor packaging or labeling
- Therapeutic failures (product didn't work)

Report SERIOUS adverse events. An event is serious when the patient outcome is:
- Death
- Life-threatening
- Hospitalization - initial or prolonged
- Disability or permanent damage
- Congenital anomaly/birth defect
- Required intervention to prevent permanent impairment or damage (devices)
- Other serious (important medical events)

Report even if:
- You’re not certain the product caused the event
- You don’t have all the details

How to report:
- Just fill in the sections that apply to your report
- Use section D for all products except medical devices
- Attach additional pages if needed
- Use a separate form for each patient
- Report either to FDA or the manufacturer (or both)

Other methods of reporting:
- 1-800-FDA-0178 - To FAX report
- 1-800-FDA-1088 - To report by phone
- www.fda.gov/medwatch/report.htm - To report online

If your report involves a serious adverse event with a device and it occurred in a facility outside a doctor's office, that facility may be legally required to report to FDA and/or the manufacturer. Please notify the person in that facility who would handle such reporting.

If your report involves a serious adverse event with a vaccine, call 1-800-822-7967 to report.

Confidentiality: The patient's identity is held in strict confidence by FDA and protected to the fullest extent of the law. FDA will not disclose the reporter's identity in response to a request from the public, pursuant to the Freedom of Information Act. The reporter's identity, including the identity of a self-reporter, may be shared with the manufacturer unless requested otherwise.

The information in this box applies only to requirements of the Paperwork Reduction Act of 1995

The burden time for this collection of information has been estimated to average 36 minutes per response, including the time to review instructions, search existing data sources, gather and maintain the data needed, and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to:

Department of Health and Human Services
Food and Drug Administration
Office of Chief Information Officer
Paperwork Reduction Act (PRA) Staff
PRASstaff@fda.hhs.gov

Please DO NOT return this form to the PRA Staff e-mail to the left.

OMB statement: "An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number."

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

FORM FDA 3500 (2/13) (Back) Please Use Address Provided Below - Fold in Thirds, Tape and Mail

DEPARTMENT OF HEALTH & HUMAN SERVICES
Public Health Service
Food and Drug Administration
Rockville, MD 20857

Official Business
Penalty for Private Use $300

BUSINESS REPLY MAIL
FIRST CLASS MAIL PERMIT NO. 946 ROCKVILLE MD
POSTAGE WILL BE PAID BY FOOD AND DRUG ADMINISTRATION

MEDWATCH
The FDA Safety Information and Adverse Event Reporting Program
Food and Drug Administration
5600 Fishers Lane
Rockville, MD 20852-9787
B.5. Describe Event or Problem (continued)

B.6. Relevant Tests/Laboratory Data, Including Dates (continued)

B.7. Other Relevant History, Including Preexisting Medical Conditions (e.g., allergies, race, pregnancy, smoking and alcohol use, hepatic/renal dysfunction, etc.) (continued)

F. Concomitant Medical Products and Therapy Dates (Exclude treatment of event) (continued)